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Rachel Sandford



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House of Commons - Science and Technology Committee: Clinical Trials - HC 104 Great Britain: Parliament: House of Commons: Science and Technology Committee, 2013-09-17 Many of the trials taking place today are unregistered and unpublished meaning that the information that they generate remains invisible to both the scientific community and the public This undermines public trust slowing the pace of medical advancement and potentially putting patients at risk All trials conducted on NHS treatments and all other trials receiving public funding should be prospectively registered and their results published in a scientific journal While the focus should be on implementing this change for future trials the Government must also do what it can to ensure that historic trials are registered and published particularly where they have been publically funded The Government should also take steps to facilitate greater sharing of the raw data generated during a trial in a responsible and controlled way with the knowledge and consent of patients The report also draws attention to the recent fall in the number of trials taking place in the UK It finds that the need for multiple governance approvals from participating NHS organisations remained the biggest barrier to setting up a UK trial but that lack of public awareness was also a key issue Recruiting participants can also be a challenge The report calls on the Government to take its recommendations into account in ongoing discussions regarding the revision of European clinical trials legislation and in its response to the European Medicines Agency's consultation on the release of clinical trial data which closes at the end of this

month The Future of Drone Use Bart Custers,2016-10-15 Given the popularity of drones and the fact that they are easy and cheap to buy it is generally expected that the ubiquity of drones will significantly increase within the next few years This raises questions as to what is technologically feasible now and in the future what is acceptable from an ethical point of view and what is allowed from a legal point of view Drone technology is to some extent already available and to some extent still in development The aim and scope of this book is to map the opportunities and threats associated with the use of drones and to discuss the ethical and legal issues of the use of drones This book provides an overview of current drone technologies and applications and of what to expect in the next few years The question of how to regulate the use of drones in the future is addressed by considering conditions and contents of future drone legislation and by analyzing issues surrounding privacy and safeguards that can be taken As such this book is valuable to scholars in several disciplines such as law ethics sociology politics and public administration as well as to practitioners and others who may be confronted with the use of drones in their work such as professionals working in the military law enforcement disaster management and infrastructure management Individuals and businesses with a specific interest in drone use may also find in the nineteen contributions contained in this volume unexpected perspectives on this new field of research and innovation Bart Custers is Associate Professor and Head of Research at eLaw the Center for Law and Digital Technologies at Leiden University The Netherlands He has presented his work at international conferences in the United States China Japan the Middle East and throughout Europe and has published over 80 scientific professional and popularizing publications including three books *MOOCs and Open Education in the Global South* Ke Zhang,Curtis Bonk,Thomas Reeves,Thomas Reynolds,2019-11-05 With e learning technologies evolving and expanding at high rates organizations and institutions around the world are integrating massive open online courses MOOCs and other open educational resources OERs MOOCs and Open Education in the Global South explores the initiatives that are leveraging these flexible systems to educate train and empower populations previously denied access to such opportunities Featuring contributors leading efforts in rapidly changing nations and regions this wide ranging collection grapples with accreditation credentialing quality standards innovative assessment learner motivation and attrition and numerous other issues The provocative narratives curated in this volume demonstrate how MOOCs and OER can be effectively designed and implemented in vastly different ways in particular settings as detailed by experts from Asia Latin America the Middle East Africa the Pacific Oceania and the Caribbean This comprehensive text is an essential resource for policy makers instructional designers practitioners administrators and other MOOC and OER community stakeholders

Pharmaceutical Freedom Jessica Flanigan,2017-07-03 Jessica Flanigan defends patients rights of self medication on the grounds that same moral reasons against medical paternalism in clinical contexts are also reasons against paternalistic pharmaceutical policies including prohibitive approval processes and prescription requirements **Intellectual Property and Public Health in the Developing World** Monirul Azam,2016-05-30 Across the world developing countries are

attempting to balance the international standards of intellectual property concerning pharmaceutical patents against the urgent need for accessible and affordable medicines In this timely and necessary book Monirul Azam examines the attempts of several developing countries to walk this fine line He evaluates the experiences of Brazil China India and South Africa for lessons to guide Bangladesh and developing nations everywhere Azam s legal expertise concern for public welfare and compelling grasp of principal case studies make Intellectual Property and Public Health in the Developing World a definitive work The developing world is striving to meet the requirements of the World Trade Organization s TRIPS Agreement on intellectual property This book sets out with lucidity and insight the background of the TRIPS Agreement and its implications for pharmaceutical patents the consequences for developing countries and the efforts of certain representative nations to comply with international stipulations while still maintaining local industry and public health Azam then brings the weight of this research to bear on the particular case of Bangladesh offering a number of specific policy recommendations for the Bangladeshi government and for governments the world over Intellectual Property and Public Health in the Developing World is a must read for public policy makers academics and students non governmental organizations and readers everywhere who are interested in making sure that developing nations meet the health care needs of their people

Parliamentary Debates (Hansard). Great Britain. Parliament. House of Commons,2014 Casarett & Doull's Toxicology: The Basic Science of Poisons, 9th Edition Curtis D. Klaassen,2018-11-23 Toxicology s gold standard text completely updated to reflect the latest breakthroughs and discoveries A Doody s Core Title for 2024 2021 Casarett Doull s Toxicology The Basic Science of Poisons Ninth Edition equips you with an unsurpassed understanding of modern toxicology including the key principles concepts mechanisms chemical specific toxicity and modes of thought that are the foundation of the discipline This trusted classic not only delivers a comprehensive review of the essential components of toxicology it offers the most up to date revealing and in depth look at the systemic responses of toxic substance available anywhere Casarett Doull s Toxicology The Basic Science of Poisons Ninth Edition is logically divided into seven sections General Principles of Toxicology Disposition of Toxicants Non Organ Directed Toxicity Target Organ Toxicity Toxic Agents Environmental Toxicology Applications of Toxicology Many new contributors capture the progress made in toxicology over the past few years This edition is markedly updated from the previous edition with more than one third of the chapters authored by scientists who have not made previous contributions to the book Sharing their expertise they deliver dynamic new coverage of the importance of apoptosis autophagy cytokines growth factors oncogenes cell cycling receptors gene regulation protective mechanisms repair mechanisms transcription factors signaling pathways transgenic mice knock out mice humanized mice polymorphisms microarray technology second generation sequencing genomics proteomics epigenetics exposome microbiota read across adverse outcome pathways high content screening computational toxicology innovative test methods and organ on a chip in understanding the mechanisms of toxicity and the regulation of chemicals A true essential If you are in need of

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